Amendments to the Claims:

This listing of claims will replace all prior versions, and listings, of claims in the application:

Listing Of Claims:

1. (Currently Amended) A drug delivery molecule comprising:

a polymerized carboxylic acid molecular scaffold having a plurality of free carboxylic acid groups;

a plurality of biologically active molecular modules, each being covalently linked to the same polymerized carboxylic acid molecular scaffold, wherein said active modules comprise: at least one targeting module for promoting cellular uptake by a target cell; and at least one pro-drug module for altering cellular metabolism of the target cell;

wherein at least one active module comprises a polypeptide and/or polynucleotide; and wherein the scaffold is a homopolymer.

- 2. (Original) The drug delivery molecule according to claim 1, wherein the pro-drug is selected to inhibit expression of tumor-specific proteins.
- 3. (Original) The drug delivery molecule according to claim 1, wherein the polymerized carboxylic acid molecular scaffold is poly (\(\beta\)-L-malic acid).
- 4. (Previously Presented) The drug delivery molecule according to claim 3, wherein the poly (B-L-malic acid) has a weight-averaged molecular weight (Mw) between 2,500 and 100,000.
- 5. (Previously Presented) The drug delivery molecule according to claim 4, wherein the poly (B-L-malic acid) has a weight-averaged molecular weight (Mw) of at least about 5,000.

- 6. (Original) The drug delivery molecule according to claim 1, wherein each molecule of the polymerized carboxylic acid molecular scaffold has at least about 50 free carboxylic acid groups.
- 7. (Original) The drug delivery molecule according to claim 1, wherein the plurality of molecular modules further includes a molecular module for promoting disruption of biomembranes.
- 8. (Original) The drug delivery molecule according to claim 7, wherein said molecular module for promoting disruption of biomembranes comprises a molecule having lipophilic characteristics and groups that are charged at physiologic pH and become uncharged at lysosomal pH thereby increasing lipophilicity of said molecular module.
- 9. (Original) The drug delivery molecule according to claim 1, wherein the plurality of active molecular modules further includes a molecular module for prolonging circulation of the drug delivery molecule.
- 10. (Original) The drug delivery molecule according to claim 9, wherein the molecular module for prolonging circulation of the drug delivery molecule comprises polyethylene glycol.
- 11. (Original) The drug delivery molecule according to claim 1, wherein the plurality of active molecular modules further includes a reporter module for determining cellular uptake of the drug delivery molecule.
- 12. (Original) The drug delivery molecule according to claim 11, wherein the reporter module comprises a fluorescent molecule.
- 13. (Original) The drug delivery molecule according to claim 1, wherein the targeting molecule is selected to promote penetration of the blood brain barrier.

Claims 14-17 (Canceled)

- 18. (Original) The drug delivery molecule according to claim 1, wherein the pro-drug molecular module is linked to the polymerized carboxylic acid molecular scaffold by a cleavable linkage that is cleaved when the drug delivery molecule enters a cell.
- 19. (Original) The drug delivery molecule according to claim 18, wherein the cleavable linkage is a disulfide linkage.
- 20. (Original) The drug delivery molecule according to claim 1, wherein the pro-drug molecular module comprises an antisense molecule.
- 21. (Original) The drug delivery molecule according to claim 20, wherein the antisense molecule is a morpholino antisense molecule.
- 22. (Original) The drug delivery molecule according to claim 20, wherein the antisense molecule interferes with production of laminin-8.
- 23. (Original) The drug delivery molecule according to claim 22, wherein the antisense molecule interferes with production of laminin-8 by altering production of a laminin subunit selected from the group consisting of α4 laminin and β1 laminin.

Claims 24-28 (Canceled)